

NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a,
 CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
 AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002
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FILE 'REGISTRY' ENTERED AT 07:13:02 ON 14 MAR 2003
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STRUCTURE FILE UPDATES: 12 MAR 2003 HIGHEST RN 498527-50-7
DICTIONARY FILE UPDATES: 12 MAR 2003 HIGHEST RN 498527-50-7

TSCA INFORMATION NOW CURRENT THROUGH MAY 20, 2002

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

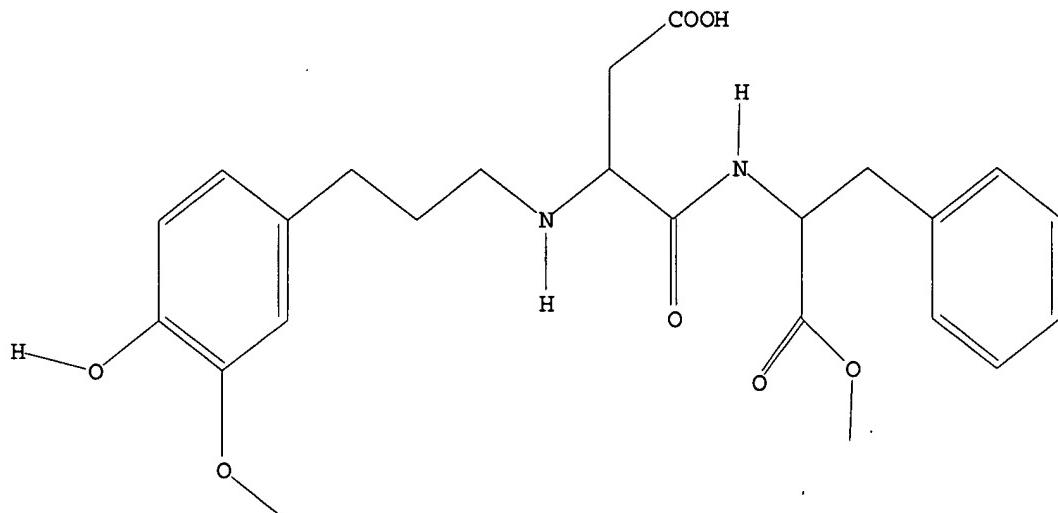
Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=>
Uploading 10091500 aspartame derivative.str

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS
L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> search l1 exact full
FULL SEARCH INITIATED 07:13:33 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 13 TO ITERATE

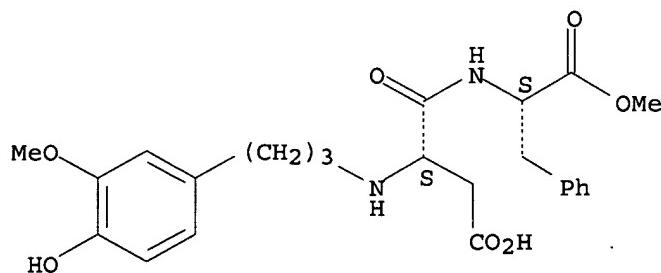
100.0% PROCESSED 13 ITERATIONS 1 ANSWERS
SEARCH TIME: 00.00.01

L2 1 SEA EXA FUL L1

=> d scan

L2 1 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Phenylalanine, N-[3-(4-hydroxy-3-methoxyphenyl)propyl]-L-.alpha.-
aspartyl-, 2-methyl ester (9CI)
MF C24 H30 N2 O7

Absolute stereochemistry.



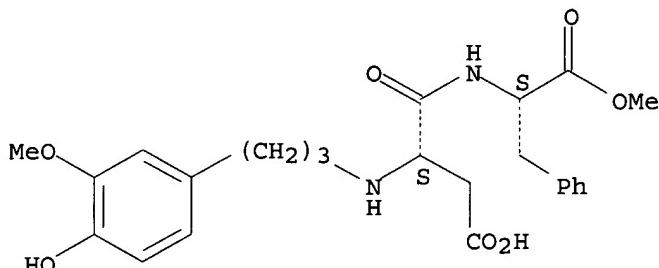
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> d 12

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS
RN 329326-75-2 REGISTRY
CN L-Phenylalanine, N-[3-(4-hydroxy-3-methoxyphenyl)propyl]-L-.alpha.-aspartyl-, 2-methyl ester (9CI) (CA INDEX NAME)
OTHER NAMES:
CN Ararame
FS STEREOSEARCH
MF C24 H30 N2 O7
SR CA
LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1962 TO DATE)
3 REFERENCES IN FILE CAPLUS (1962 TO DATE)

=> file caplus	SINCE FILE	TOTAL
COST IN U.S. DOLLARS	ENTRY	SESSION
FULL ESTIMATED COST	51.83	52.04

FILE 'CAPLUS' ENTERED AT 07:13:55 ON 14 MAR 2003
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FILE COVERS 1907 - 14 Mar 2003 VOL 138 ISS 12
FILE LAST UPDATED: 13 Mar 2003 (20030313/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> l2
L3 3 L2

=> d l3 1-3 ti fbib abs

L3 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2003 ACS
TI Gustatory responses of pigs to sixty compounds tasting sweet to humans
AN 2002:351144 CAPLUS
DN 137:336847
TI Gustatory responses of pigs to sixty compounds tasting sweet to humans
AU Nofre, C.; Glaser, D.; Tinti, J.-M.; Wanner, M.
CS Faculty of Medicine of Lyon Laennec, University of Lyon, Lyon, Fr.
SO Journal of Animal Physiology and Animal Nutrition (2002), 86(3-4), 90-96
CODEN: JAPNEF; ISSN: 0931-2439
PB Blackwell Wissenschafts-Verlag GmbH
DT Journal
LA English
AB The gustatory responses of pigs to 60 compds. perceived as sweet by humans

were studied via a semi-quant. behavioral method derived from the Richter two-bottle preference test. Among the 60 compds. tested 35 are effective in pigs, but with an effectiveness much lower in pigs than in humans. Lugduname and carrelame, which are the two most potent sweeteners in humans, are also the most effective compds. in pigs.

RE.CNT 31 THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS
TI Process for producing aspartyl dipeptide ester derivatives
AN 2001:833348 CAPLUS
DN 135:358168
TI Process for producing aspartyl dipeptide ester derivatives
IN Kawahara, Shigeru; Nagashima, Kazutaka; Takemoto, Tadashi
PA Ajinomoto Co., Inc., Japan
SO PCT Int. Appl., 25 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
	-----	-----	-----	-----	-----	
PI	WO 2001085761	A1	20011115	WO 2001-JP3479	20010423	
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,				

BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
 JP 2000-137028 A 20000510
 EP 1283213 A1 20030212 EP 2001-922023 20010423
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
 JP 2000-137028 A 20000510
 WO 2001-JP3479 W 20010423

OS CASREACT 135:358168; MARPAT 135:358168

AB This document discloses a process for conveniently producing on an industrial scale in high yield

N- [N- [3- (phenyl) propyl] -L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester derivs., which are expected to be sweeteners, by reductively alkylating aspartame with 3-phenyl-2-propenyl aldehyde derivs. under hydrogen in the presence of a catalyst and a base.

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS
 TI Process for producing and purifying aspartame derivative as sweetener
 AN 2001:185780 CAPLUS
 DN 134:223039
 TI Process for producing and purifying aspartame derivative as sweetener
 IN Amino, Yusuke; Yuzawa, Kazuko; Takemoto, Tadashi
 PA Ajinomoto Co., Inc., Japan
 SO PCT Int. Appl., 39 pp.
 CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001018034	A1	20010315	WO 2000-JP5665	20000823
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	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 1999-253498 A	19990907
AU	2000067273	A5	20010410	AU 2000-67273	20000823
				JP 1999-253498 A	19990907
				WO 2000-JP5665 W	20000823
US	2002147361	A1	20021010	US 2002-91500	20020307
				JP 1999-253498 A	19990907
				WO 2000-JP5665 A120000823	

OS CASREACT 134:223039

AB This document discloses the following : a method for industrially producing

N- [N- [3- (3-methoxy-4-hydroxyphenyl) propyl] -L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester which is useful as a sweetener, in particular, a process for producing the target compd. in a high yield by the reductive alkylation reaction of aspartame with 3-(3-methoxy-4-hydroxyphenyl)propionaldehyde or its deriv.; a method of effectively purifying the target compd. contaminated with impurities invading thereinto at various prodn. stages (involving methods other than the above-described reductive alkylation), more particularly, a method of sepg. the target compd. in the form of highly pure crystals; the crystals;

sweeteners contg. the same; and utilization thereof in various products which are to be sweetened.

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> file reg		SINCE FILE	TOTAL
COST IN U.S. DOLLARS		ENTRY	SESSION
FULL ESTIMATED COST		8.91	60.95
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)		SINCE FILE	TOTAL
CA SUBSCRIBER PRICE		ENTRY	SESSION
		-1.95	-1.95

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STRUCTURE FILE UPDATES: 12 MAR 2003 HIGHEST RN 498527-50-7
DICTIONARY FILE UPDATES: 12 MAR 2003 HIGHEST RN 498527-50-7

TSCA INFORMATION NOW CURRENT THROUGH MAY 20, 2002

Please note that search-term pricing does apply when conducting SmartSELECT searches.

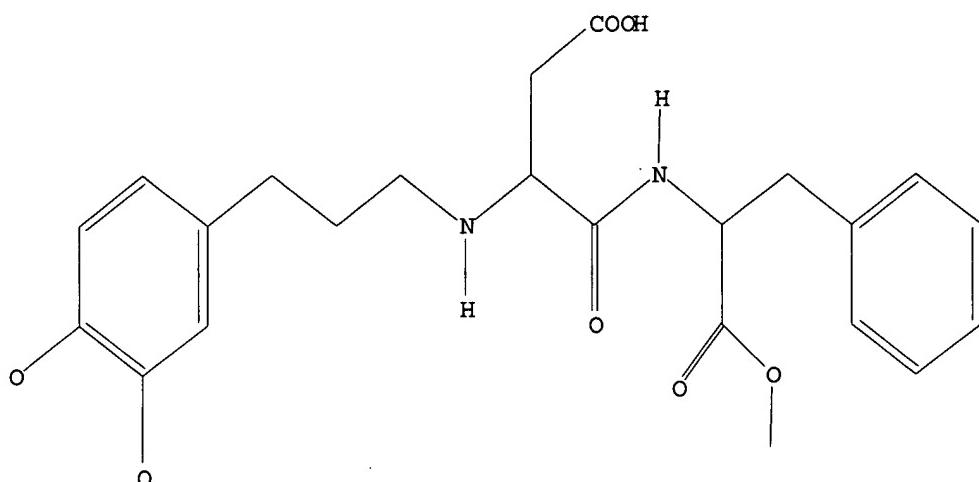
Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=>
Uploading 10091500 aspartame derivative sss.str

L4 STRUCTURE uploaded

=> d 14
L4 HAS NO ANSWERS
L4 STR



Structure attributes must be viewed using STN Express query preparation.

```
=> search 14 sss full
FULL SEARCH INITIATED 07:17:08 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 196 TO ITERATE
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100.0% PROCESSED 196 ITERATIONS
SEARCH TIME: 00.00.01

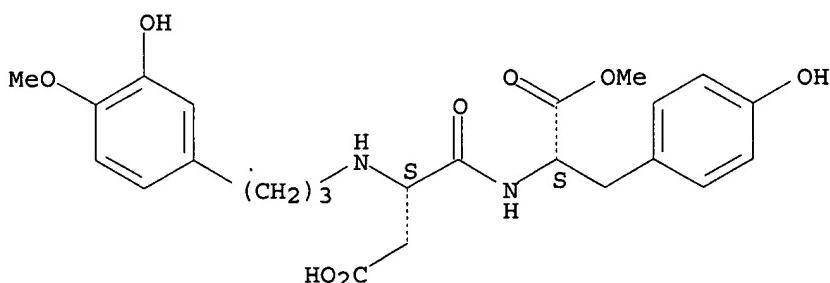
20 ANSWERS

L5 20 SEA SSS FUL L4

=> d scan

```
L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Tyrosine, N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl-,
2-methyl ester (9CI)
MF C24 H30 N2 O8
```

Absolute stereochemistry.

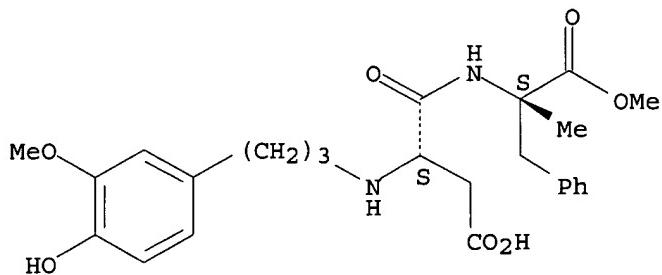


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):20

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Phenylalanine, N-[3-(4-hydroxy-3-methoxyphenyl)propyl]-L-.alpha.-
aspartyl-.alpha.-methyl-, 2-methyl ester (9CI)
MF C25 H32 N2 O7

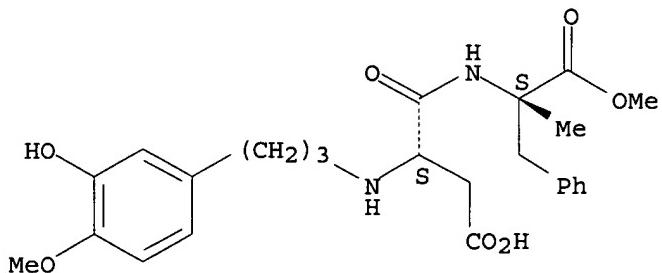
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Phenylalanine, N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-
aspartyl-.alpha.-methyl-, 2-methyl ester (9CI)
MF C25 H32 N2 O7

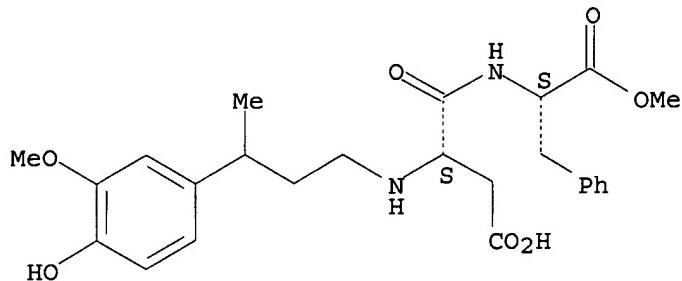
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Phenylalanine,
N-[3-(4-hydroxy-3-methoxyphenyl)butyl]-L-.alpha.-aspartyl-
, 2-methyl ester (9CI)
MF C25 H32 N2 O7

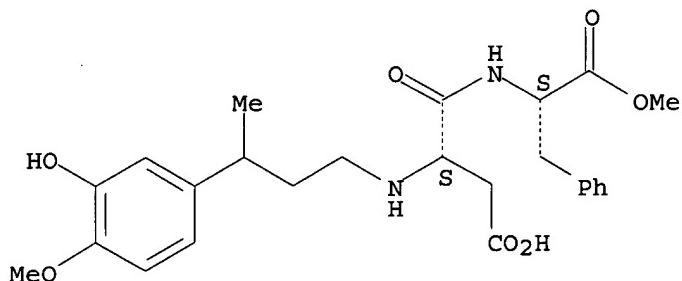
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Phenylalanine,
N- [3- (3-hydroxy-4-methoxyphenyl)butyl]-L-.alpha.-aspartyl-
, 2-methyl ester (9CI)
MF C25 H32 N2 O7

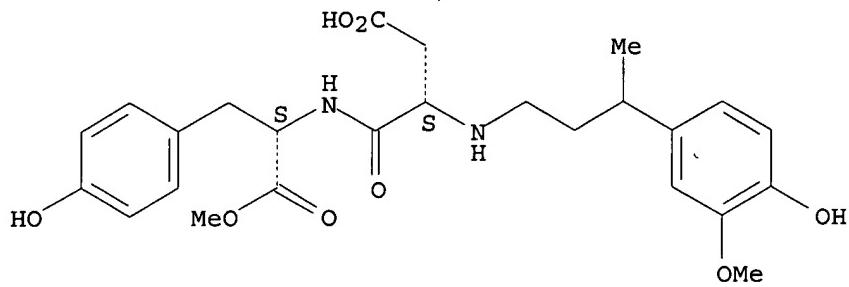
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
IN L-Tyrosine, N- [3- (4-hydroxy-3-methoxyphenyl)butyl]-L-.alpha.-aspartyl-,
2-methyl ester (9CI)
MF C25 H32 N2 O8

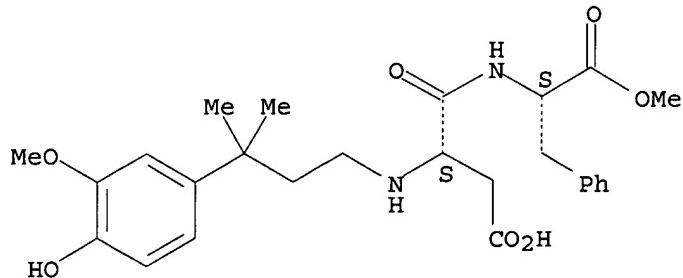
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
 N- [3 - (4-hydroxy-3-methoxyphenyl) -3-methylbutyl] -L-.alpha.-
 aspartyl-, 2-methyl ester (9CI)
 MF C26 H34 N2 O7

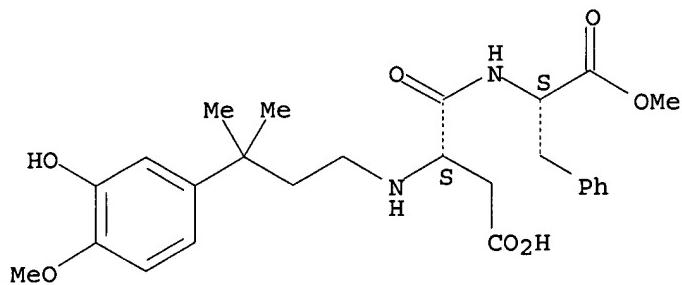
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
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 aspartyl-, 2-methyl ester (9CI)
 MF C26 H34 N2 O7

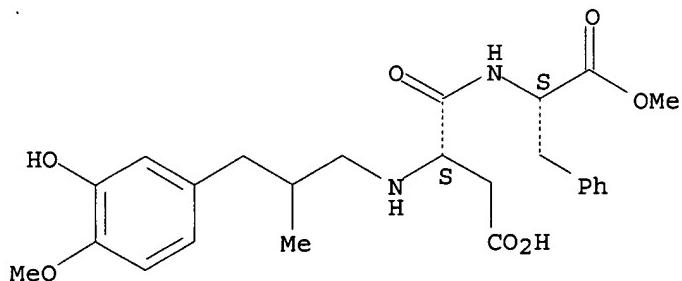
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(3-hydroxy-4-methoxyphenyl)-2-methylpropyl]-L-alpha.-aspartyl-, 2-methyl ester (9CI)
 MF C25 H32 N2 O7

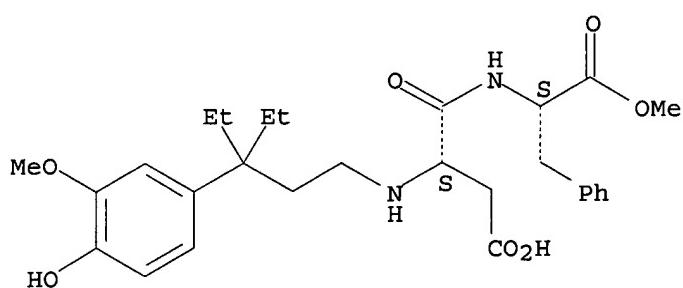
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
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 aspartyl-, 2-methyl ester (9CI)
 MF C28 H38 N2 O7

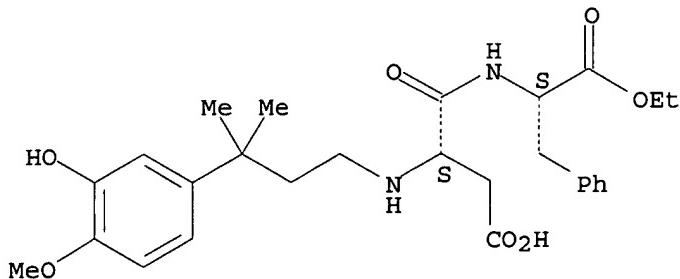
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
 N-[3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyl]-L-.alpha.-
 aspartyl-, 2-ethyl ester (9CI)
 MF C27 H36 N2 O7

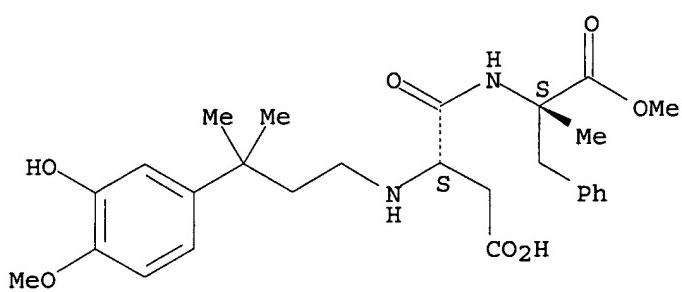
Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
 N-[3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyl]-L-.alpha.-
 aspartyl-.alpha.-methyl-, 2-methyl ester (9CI)
 MF C27 H36 N2 O7

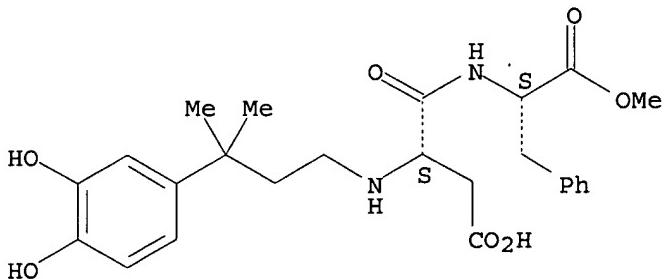
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(3,4-dihydroxyphenyl)-3-methylbutyl]-L-.alpha.-aspartyl-, 2-methyl ester (9CI)
 MF C25 H32 N2 O7

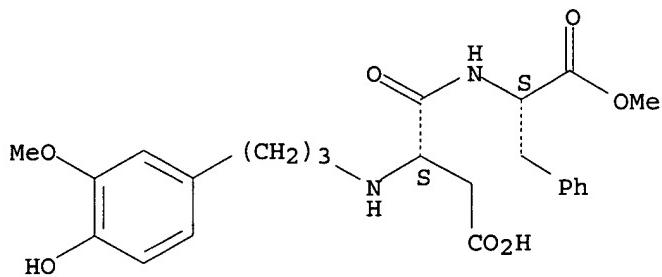
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(4-hydroxy-3-methoxyphenyl)propyl]-L-.alpha.-aspartyl-, 2-methyl ester (9CI)
 MF C24 H30 N2 O7

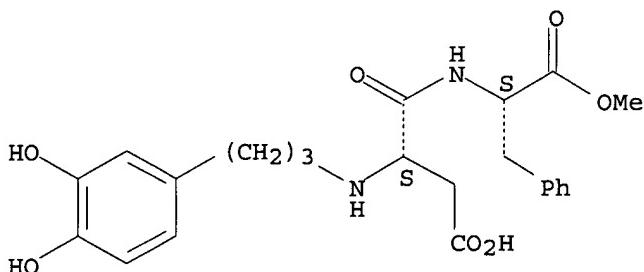
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(3,4-dihydroxyphenyl)propyl]-L-.alpha.-aspartyl-,
 2-methyl ester (9CI)
 MF C23 H28 N2 O7

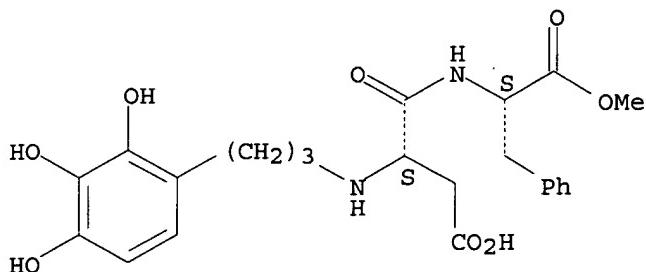
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
 N-[3-(2,3,4-trihydroxyphenyl)propyl]-L-.alpha.-aspartyl-,
 2-methyl ester (9CI)
 MF C23 H28 N2 O8

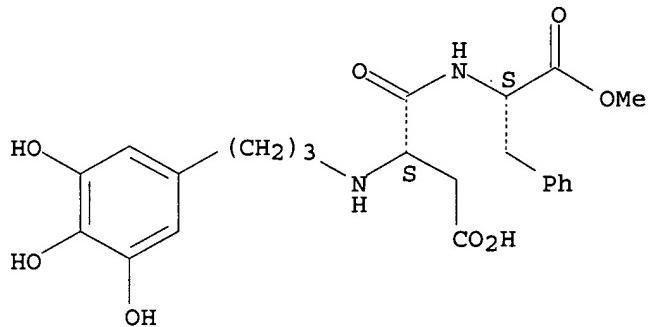
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine,
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 2-methyl ester (9CI)
 MF C23 H28 N2 O8

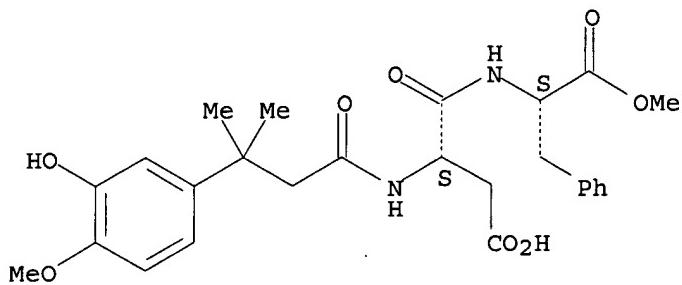
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(3-hydroxy-4-methoxyphenyl)-3-methyl-1-oxobutyl]-L-.alpha.-aspartyl-, 2-methyl ester (9CI)
 MF C26 H32 N2 O8

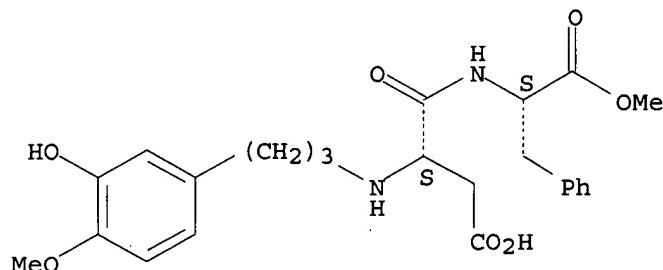
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl-, 2-methyl ester (9CI)
 MF C24 H30 N2 O7

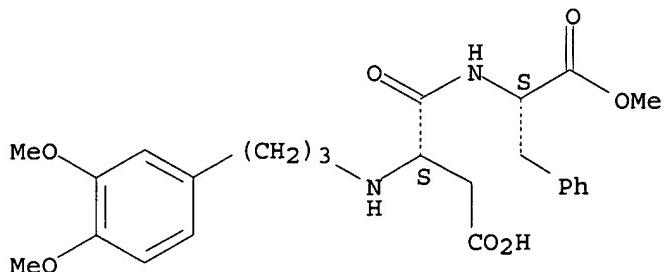
Absolute stereochemistry.



***PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

L5 20 ANSWERS REGISTRY COPYRIGHT 2003 ACS
 IN L-Phenylalanine, N-[3-(3,4-dimethoxyphenyl)propyl]-L-.alpha.-aspartyl-, 2-methyl ester (9CI)
 MF C25 H32 N2 O7

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> file caplus			
COST IN U.S. DOLLARS	SINCE FILE	TOTAL	
	ENTRY	SESSION	
FULL ESTIMATED COST	148.55	209.50	
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL	
	ENTRY	SESSION	
CA SUBSCRIBER PRICE	0.00	-1.95	

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FILE COVERS 1907 - 14 Mar 2003 VOL 138 ISS 12
 FILE LAST UPDATED: 13 Mar 2003 (20030313/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> 15
 L6 14 L5

=> d his

(FILE 'HOME' ENTERED AT 07:12:53 ON 14 MAR 2003)

FILE 'REGISTRY' ENTERED AT 07:13:02 ON 14 MAR 2003

L1 STRUCTURE UPLOADED
L2 1 SEARCH L1 EXACT FULL

FILE 'CAPLUS' ENTERED AT 07:13:55 ON 14 MAR 2003
L3 3 L2

FILE 'REGISTRY' ENTERED AT 07:16:35 ON 14 MAR 2003
L4 STRUCTURE UPLOADED
L5 20 SEARCH L4 SSS FULL

FILE 'CAPLUS' ENTERED AT 07:17:35 ON 14 MAR 2003
L6 14 L5

=> 16 not 13
L7 11 L6 NOT L3

=> d 17 1-11 ti

L7 ANSWER 1 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Process for producing cinnamyl aldehyde derivatives and use thereof as intermediate for aspartame derivative

L7 ANSWER 2 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Aspartame derivative crystals

L7 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Preparation of 3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyric acid derivative as novel intermediate for sweetener with high sweetness and process for producing the same

L7 ANSWER 4 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Process for producing N-(3-methyl-3-phenylbutyl)aspartame derivative, crystals thereof, novel production of aldehyde intermediates therefor and process for producing the intermediate

L7 ANSWER 5 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Sweetener compositions with high degree of sweetness having improved sweetness, supplements and utilization thereof

L7 ANSWER 6 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Sweetener compositions with high degree of sweetness having improved sweetness, supplements and utilization thereof

L7 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Process for the production of aspartyldipeptide ester derivatives, novel intermediates therefor and process for the production of the intermediates

L7 ANSWER 8 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Preparation of aspartyl dipeptides and their use as sweeteners

L7 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI N-Alkylaspartyldipeptide ester derivatives and sweeteners

L7 ANSWER 10 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Novel aspartyl dipeptide ester derivatives as sweeteners

L7 ANSWER 11 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Preparation of aspartyl dipeptide ester derivatives as sweeteners

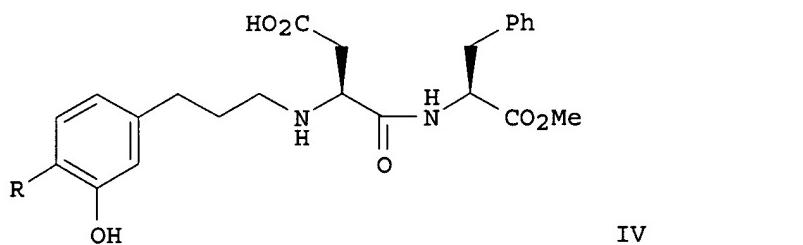
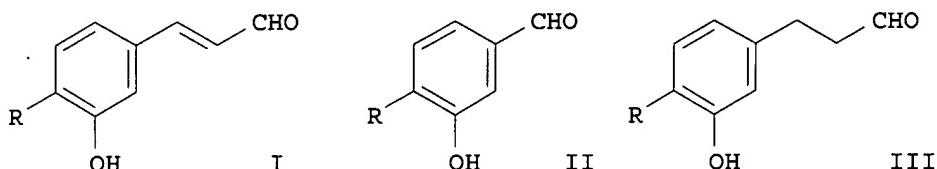
=> d 17 1-11 ti fbib abs

L7 ANSWER 1 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Process for producing cinnamyl aldehyde derivatives and use thereof as intermediate for aspartame derivative
AN 2001:851092 CAPLUS
DN 135:371997
TI Process for producing cinnamyl aldehyde derivatives and use thereof as intermediate for aspartame derivative
IN Mori, Kenichi; Fujita, Shinji; Funakoshi, Nao; Takemoto, Tadashi
PA Ajinomoto Co., Inc., Japan
SO PCT Int. Appl., 29 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001087813	A1	20011122	WO 2001-JP3545	20010424
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 2000-142811 A	20000516
EP 1283197	A1	20030212	EP 2001-922073	20010424
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR		JP 2000-142811 A	20000516
			WO 2001-JP3545	W 20010424

OS CASREACT 135:371997; MARPAT 135:371997

GI



AB Described is an industrial process for conveniently and efficiently

producing highly pure cinnamyl aldehyde derivs. (I; R = H, C1-4 alkyl or alkoxy) such as (2E)-(3-hydroxy-4-methoxy)cinnamyl aldehyde which comprises reacting a benzaldehyde deriv. (II; R = same as above) (for example, isovanillin) with acetaldehyde in the presence of an alkali, preferably adding acetaldehyde in portions in an aq. soln. at a low temp. The cinnamyl aldehyde derivs. (I) thus obtained are selectively reduced into 3-(3-hydroxy-4-substituted phenyl)propionaldehydes (III; R = same as above). These compds. III are further subjected to reductive alkylation with aspartame to efficiently give N-[N-[3-(3-hydroxy-4-substituted phenyl)propyl]-L-.alpha.-aspartyl]-L-phenylalanine 1-Me esters (IV; R =

H, C1-4 alkyl or alkoxy), which are useful as sweeteners with high sweetness.

Thus, 121.72 g isovanillin and 320 g NaOH were dissolved in 2,000 mL H₂O and cooled to -10.degree., followed by continuously adding 290 g 28 wt.% aq. acetaldehyde over a period of 45 h, and the resulting mixt. was stirred for 1 h, treated with 768.1 g 36 wt.% aq. HCl, and filtered to give 324 g cryst. product. The latter product was dispersed in 500 mL

H₂O at 25.degree., treated with 97.5 g 25 wt.% aq. NaOH for dissoln., stirred with 4 g activated charcoal and 16 g celite, and filtered. The filtrate was neutralized with 55.4 g 36 wt.% aq. HCl to give 185.5 g cryst.

product

which was vacuum-dried, dispersed in 275 mL MeOH at 60.degree., stirred for 2 h, cooled to room temp., and filtered to give, after drying the wet crystals, 83.2 g (2E)-3-hydroxy-4-methoxycinnamaldehyde (98% purity) in 57% yield. The latter compd. (5.00 g) and 300 mg 5% Pd-Al₂O₃ were added to 80 mL MeOH and stirred under H atm. at 35.degree. for 24 h, followed

by filtration for removal of the catalyst and washing the catalyst with 10 mL

MeOH, to give a MeOH soln. of

3-(3-hydroxy-4-methoxyphenyl)propionaldehyde (87% yield). The latter soln. (8.15 g) contg. 1.50 g of the aldehyde and 2.57 g aspartame were added to a 4:1 mixt. of MeOH and H₂O, followed by adding 0.7 g 10% Pd-C contg. 50% H₂O, and the resulting mixt. was stirred at 35.degree. under H atm. for 48 h to give 71% N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester.

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Aspartame derivative crystals
AN 2001:489419 CAPLUS
DN 135:60486
TI Aspartame derivative crystals
IN Nagashima, Kazutaka; Aoki, Yuichi; Ono, Eriko; Takemoto, Tadashi
PA Ajinomoto Co., Inc., Japan
SO PCT Int. Appl., 29 pp.
CODEN: PIXXD2

DT Patent.

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001047949	A1	20010705	WO 2000-JP9247	20001225
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,				

SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
 YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
 DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
 BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
 JP 1999-373257 A 19991228
 AU 2001022259 A5 20010709 AU 2001-22259 20001225
 JP 1999-373257 A 19991228
 WO 2000-JP9247 W 20001225
 EP 1245573 A1 20021002 EP 2000-985895 20001225
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
 JP 1999-373257 A 19991228
 WO 2000-JP9247 W 20001225
 BR 2000016316 A 20021203 BR 2000-16316 20001225
 JP 1999-373257 A 19991228
 WO 2000-JP9247 W 20001225
 US 2003009050 A1 20030109 US 2002-183652 20020628
 JP 1999-373257 A 19991228
 WO 2000-JP9247 A120001225

AB Com. favorable crystals of N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl]-L-phenylalanine-Me ester (I) were given. Compared to amorphous aspartame, I have better stability, and higher purity and sweetness. Physicochem. characteristics of the I crystals were also given.

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

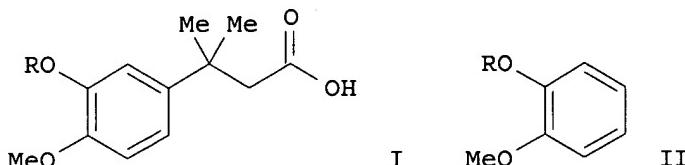
L7 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2003 ACS
 TI Preparation of 3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyric acid derivative as novel intermediate for sweetener with high sweetness and process for producing the same
 AN 2001:396837 CAPLUS
 DN 135:5819
 TI Preparation of 3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyric acid derivative as novel intermediate for sweetener with high sweetness and process for producing the same
 IN Kawahara, Shigeru; Mori, Kenichi; Nagashima, Kazutaka; Takemoto, Tadashi
 PA Ajinomoto Co., Inc., Japan
 SO PCT Int. Appl., 26 pp.
 CODEN: PIXXD2
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001038297	A1	20010531	WO 2000-JP7913	20001109
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 1999-328100 A 19991118		
	AU 2001013052	A5	20010604	AU 2001-13052	20001109
				JP 1999-328100 A 19991118	
				WO 2000-JP7913 W 20001109	

EP 1236713 A1 20020904 EP 2000-974890 20001109
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, MC, IE, SI,
LT, LV, FI, RO, MK, CY, AL

JP 1999-328100 A 19991118
WO 2000-JP7913 W 20001109

OS CASREACT 135:5819; MARPAT 135:5819
GI



AB The title compds. (I; R = sulfonyl-type protecting group) can be obtained by substituting the substituent at the 3-position of the benzene ring of a

butyric acid deriv. which can be easily and efficiently produced by reacting a hydroxyl-protected 2-methoxyphenol (II; R = same as above), wherein the hydroxyl group of 2-methoxyphenol is protected in the form of a sulfonate, with 3-methylcrotonic acid in the presence of an acid. By further converting the carboxyl group into a formyl group, 3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyraldehyde can be easily produced. This aldehyde deriv. can be easily derived into a compd.,

which

is excellent as a sweetener with a high sweetness, by reductive alkylation

with aspartame. Thus, 104 g AlCl₃ was added to a soln. of 240 g 2-methanesulfonyloxyanisole and 39 g 3-methylcrotonic acid, stirred at 70.degree. for 5 h and 100.degree. for 2 h, cooled to room temp., treated with 390 mL 6 N HCl, stirred vigorously for 3 h, and extd. with 300 mL CH₂Cl₂. The org. layer was extd. with 400 mL 2 N NaOH and the sepd. aq. layer was acidified with 6 N HCl, and extd. twice with 300 mL CH₂Cl₂.

The

org. layer was concd. under reduced pressure to give a residue contg. 3-(3-methanesulfonyloxy-4-methoxyphenyl)-3-methylbutanoic acid which was treated with 300 mL 6 N NaOH, stirred at 100.degree. for 4 h, cooled to room temp., acidified with 6 N HCl, and extd. with EtOAc to give, after evapn. of the solvent from the ext. and recrystn. from toluene, 37.9% 3-(3-hydroxy-4-methoxyphenyl)-3-methylbutanoic acid (III). III (13.6 g), 22.8 g pivalic acid anhydride, and 100 mL acetone were enclosed in a high pressure hydrogenation app., purged by bubbling N for 30 min, treated

with

a soln. of 137 mg Pd(OAc)₂ and 930 mg tri(p-tolyl)phosphine in 5 mL THF, and stirred at 80.degree. under 5 MPa hydrogen pressure to give, after evapn. of acetone and column chromatog., 80%

3-(3-hydroxy-4-methoxyphenyl)-

3-methylbutyraldehyde (IV). Aspartame (8.45 g) was added to a soln. of 6.68 g IV in 272 mL 80% aq. methanol and the resulting slurry was hydrogenated in the presence of 2.86 g 10% Pd-C (50% water content) at 25.degree. for 24 h, filtered, and the filtrated was treated with 190 mL water and extd. with 250 mL PhMe. The sepd. methanol-water layer was concd. under reduced pressure to .apprx.1/2 wt., cooled from 75.degree.

to

5.degree., and filtered to collect the pptd. crystals to give, after

crystn. from 50% aq. MeOH, 67.6% N-[N-[3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyl]-L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester (98% purity), which is a sweetening agent with high sweetness (no data).

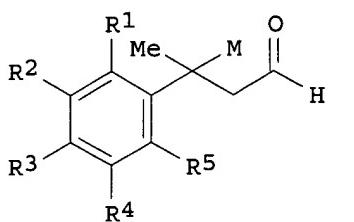
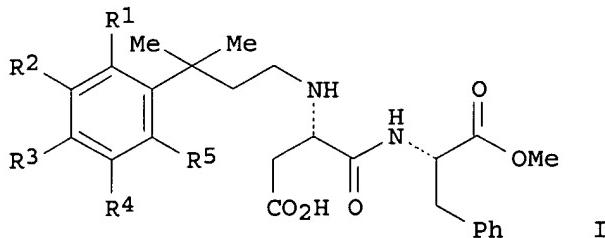
RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 4 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Process for producing N-(3-methyl-3-phenylbutyl)aspartame derivative, crystals thereof, novel production of aldehyde intermediates therefor and process for producing the intermediate
AN 2001:283984 CAPLUS
DN 134:296101
TI Process for producing N-(3-methyl-3-phenylbutyl)aspartame derivative, crystals thereof, novel production of aldehyde intermediates therefor and process for producing the intermediate
IN Kawahara, Shigeru; Nagashima, Kazutaka; Mori, Kenichi; Takemoto, Tadashi; Ono, Eriko
PA Ajinomoto Co., Inc., Japan
SO PCT Int. Appl., 50 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001027142	A1	20010419	WO 2000-JP6933	20001004
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			JP 1999-288207 A	19991008
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			JP 1999-288208 A	19991008
				JP 1999-294409 A	19991015
				JP 1999-328099 A	19991118
AU	2000075567	A5	20010423	AU 2000-75567	20001004
				JP 1999-288207 A	19991008
				JP 1999-288208 A	19991008
				JP 1999-294409 A	19991015
				JP 1999-328099 A	19991118
				WO 2000-JP6933 W	20001004
EP	1219633	A1	20020703	EP 2000-964672	20001004
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL			JP 1999-288207 A	19991008
				JP 1999-288208 A	19991008
				JP 1999-294409 A	19991015
				JP 1999-328099 A	19991118
				WO 2000-JP6933 W	20001004
BR	2000014565	A	20021119	BR 2000-14565	20001004
				JP 1999-288207 A	19991008
				JP 1999-288208 A	19991008
				JP 1999-294409 A	19991015
				JP 1999-328099 A	19991118
				WO 2000-JP6933 W	20001004
US	2002132032	A1	20020919	US 2002-117205	20020408

JP 1999-288207 A 19991008
JP 1999-288208 A 19991008
JP 1999-294409 A 19991015
JP 1999-328099 A 19991118
WO 2000-JP6933 A120001004

OS CASREACT 134:296101; MARPAT 134:296101
GI



AB The title compds. (I; R1-R5 = H, OH, C1-3 alkoxy, C1-3 alkyl, C2-3 hydroxyalkoxy; or R1 and R2 or R2 and R3 together represent methylenedioxy) are prep'd. by reductive alkylation of aspartame with 3-methyl-3-phenylbutyraldehyde derivs. (II; R1-R5 = H, OH, C1-3 alkoxy, C1-3 alkyl, benzyloxy, C2-3 hydroxyalkoxy; or R1 and R2 or R2 and R3 together represent methylenedioxy). This process is industrially advantageous since these compds. are readily crystd. and sep'd. in high purity from products contg. impurities. These compds. are useful as sweeteners having high degree of sweetness for food or beverages (no data). Thus, 3-(3-Hydroxy-4-methoxyphenyl)-3-methylbutyraldehyde (III) was prep'd. by treatment of 3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyric acid (IV) with pivalic anhydride in acetone for 30 min and hydrogenation in the presence of Pd(OAc)₂ and tri(p-tolyl)phosphine under hydrogen pressure of 5 MPa at 80.degree. for 24 h. IV was obtained by reacting 2-bromoanisole with 3-methylcrotonic acid in the presence of AlCl₃ at 70.degree. for 5 h and hydroxylation of the resulting 3-(3-bromo-4-methoxyphenyl)-3-methylbutyric acid with NaOH in the presence of CuSO₄.5H₂O in distd. water at room temp. for 1 h and 160.degree. for 10 h.

Aspartame and III were added to 80% aq. MeOH, stirred at 40.degree., hydrogenated over 10% Pd-C under hydrogen atm. at 25.degree. for 24 h, and filtered for removing the catalyst, followed by washing the catalyst with methanol, the combined filtrate was treated with water, and extd. with toluene. The sep'd. aq. methanol phase was concd. under reduced pressure to .apprx.1/2 wt., and cooled from 75.degree. to 5.degree. for crystn. The sep'd. crystals were recrystd. from 50% aq. MeOH to give 67.6% N-[N- [3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyl]-L-<α-aspartyl]-L-phenylalanine 1-Me ester as white crystals.

RE.CNT 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 5 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Sweetener compositions with high degree of sweetness having improved
sweetness, supplements and utilization thereof
AN 2001:265445 CAPLUS
DN 134:265559
TI Sweetener compositions with high degree of sweetness having improved
sweetness, supplements and utilization thereof
IN Ishii, Shoichi
PA Ajinomoto Co., Inc., Japan
SO PCT Int. Appl., 50 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI	WO 2001025263	A1	20010412	WO 2000-JP6629	20000926	
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 1999-284344 A	19991005		
				JP 1999-284345 A	19991005	
JP	2001103925	A2	20010417	JP 1999-284344	19991005	
JP	2001103926	A2	20010417	JP 1999-284345	19991005	
AU	2000073222	A5	20010510	AU 2000-73222	20000926	
				JP 1999-284344 A	19991005	
				JP 1999-284345 A	19991005	
				WO 2000-JP6629 W	20000926	
EP	1223175	A1	20020717	EP 2000-961240	20000926	
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL			JP 1999-284344 A	19991005	
				JP 1999-284345 A	19991005	
				WO 2000-JP6629 W	20000926	
BR	2000014492	A	20020820	BR 2000-14492	20000926	
				JP 1999-284344 A	19991005	
				JP 1999-284345 A	19991005	
				WO 2000-JP6629 W	20000926	
US	2003044502	A1	20030306	US 2002-115937	20020405	
				JP 1999-284344 A	19991005	
				JP 1999-284345 A	19991005	
				WO 2000-JP6629 A120000926		
OS	MARPAT 134:265559					
AB	Sweetener compns. similar to sucrose are obtained by blending aspartyl dipeptide ester derivs. (I, Markush structure claimed) such as N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl]-L- phenylalanine 1-Me ester with at least one compd. selected from the group comprising saccharides and sugar alcs., in the form of solns. These derivs. I are added to improve the taste of beverages.					
RE.CNT 4	THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT					

L7 ANSWER 6 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Sweetener compositions with high degree of sweetness having improved
sweetness, supplements and utilization thereof

AN 2001:265444 CAPLUS

DN 134:265558

TI Sweetener compositions with high degree of sweetness having improved
sweetness, supplements and utilization thereof

IN Ishii, Shoichi

PA Ajinomoto Co., Inc., Japan

SO PCT Int. Appl., 86 pp.

CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001025262	A1	20010412	WO 2000-JP6628	20000926
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 1999-283505 A 19991004 JP 1999-283506 A 19991004 JP 1999-284346 A 19991005		
	AU 2000073221	A5	20010510	AU 2000-73221	20000926
				JP 1999-283505 A 19991004 JP 1999-283506 A 19991004 JP 1999-284346 A 19991005 WO 2000-JP6628 W 20000926	
EP	1221448	A1	20020710	EP 2000-961239	20000926
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL			JP 1999-283505 A 19991004 JP 1999-283506 A 19991004 JP 1999-284346 A 19991005 WO 2000-JP6628 W 20000926	
	BR 2000014454	A	20020820	BR 2000-14454	20000926
				JP 1999-283505 A 19991004 JP 1999-283506 A 19991004 JP 1999-284346 A 19991005 WO 2000-JP6628 W 20000926	

OS MARPAT 134:265558

AB Sweetener compns. similar to sucrose are obtained by blending aspartyl
dipeptide ester derivs. (I, Markush structure claimed) such as
N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-<alpha;-aspartyl]-L-phenylalanine
1-Me ester with at least one compd. selected from the group comprising
aspartame, saccharides, sugar alcs. and oligosaccharides, so as to
enhance

the taste of I. These derivs. I are added to improve the taste of
beverages and pharmaceuticals.

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2003 ACS

TI Process for the production of aspartyldipeptide ester derivatives, novel intermediates therefor and process for the production of the intermediates

AN 2001:265443 CAPLUS

DN 134:281142

TI Process for the production of aspartyldipeptide ester derivatives, novel intermediates therefor and process for the production of the intermediates

IN Nagashima, Kazutaka; Aoki, Yuichi; Takemoto, Tadashi; Amino, Yusuke; Funakoshi, Nao; Ono, Eriko

PA Ajinomoto Co., Inc., Japan

SO PCT Int. Appl., 39 pp.

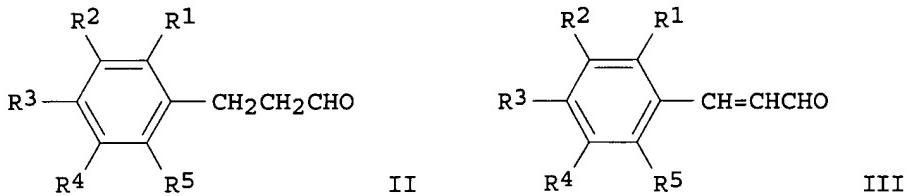
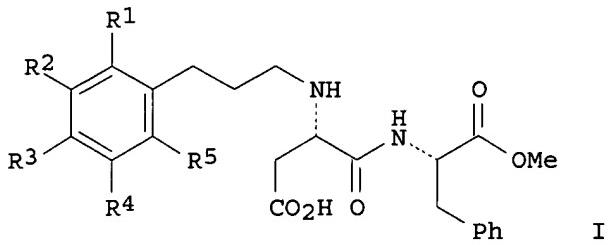
CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001025260	A1	20010412	WO 2000-JP6626	20000926
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 1999-287398 A 19991007 JP 1999-371284 A 19991227		
	AU 2000073219	A5	20010510	AU 2000-73219	20000926
				JP 1999-287398 A 19991007 JP 1999-371284 A 19991227	
				WO 2000-JP6626 W 20000926	
EP	1231215	A1	20020814	EP 2000-961237	20000926
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL			JP 1999-287398 A 19991007 JP 1999-371284 A 19991227	
				WO 2000-JP6626 W 20000926	
	US 2002133037	A1	20020919	US 2002-117196	20020408
				JP 1999-287398 A 19991007 JP 1999-371284 A 19991227	
				WO 2000-JP6626 A120000926	
OS	CASREACT 134:281142; MARPAT 134:281142				
GI					



AB Industrial and efficient processes for producing aspartylidipeptide ester derivs. of general formula (I; R1-R5 = H, OH, C1-3 alkoxy, C1-3 alkyl, benzyloxy, C2-3 hydroxyalkyloxy; or R1 and R2 or R2 and R3 together represents methylenedioxy), which are expected to serve as sweetener (no data), comprise reductive alkylation of aspartame with propionaldehydes
or cinnamaldehydes of general formulas (II) and (III) in the presence of a catalyst. Particularly, described are an industrial and efficient process for producing N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-aspartyl]-L-phenylalanine 1-Me ester (IV) which is excellent as high sweetener; useful and advantageous intermediates for the process; and efficient processes for producing the intermediates. Thus, 5.89 g aspartame and 3.42 g 3-(3-hydroxy-4-methoxyphenyl)propionaldehyde (prepn. given) were added to 200 mL 80% aq. methanol, stirred at 40.degree. for a while, and hydrogenated in the presence of 1.78 10% Pd-C at 0.1 M Pa and 40.degree. for 40 h to give 78.9% IV.

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

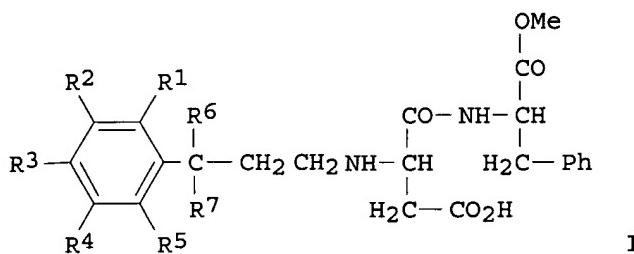
L7 ANSWER 8 OF 11 CAPLUS COPYRIGHT 2003 ACS
 TI Preparation of aspartyl dipeptides and their use as sweeteners
 AN 2001:252943 CAPLUS
 DN 134:266568
 TI Preparation of aspartyl dipeptides and their use as sweeteners
 IN Amino, Yusuke; Takemoto, Tadashi; Yuzawa, Kazuko; Nakamura, Ryoichiro
 PA Ajinomoto Co., Inc., Japan
 SO Jpn. Kokai Tokkyo Koho, 7 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2001097998	A2	20010410	JP 1999-281920	19991001
	WO 2001025261	A1	20010412	WO 2000-JP6627	20000926
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CM				

CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
 HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
 LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD,
 SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU,
 ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
 DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
 CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

JP 1999-281920 A 19991001

OS MARPAT 134:266568
GI



AB Title dipeptides I (R1-R5 = H, OH; .gtoreq.2 of R1-R5 = OH; R6, R7 = H, C1-3 alkyl) or their salts are prepd. Thus, .beta.-O-benzyl-L-.alpha.-aspartyl-L-phenylalanine Me ester was treated with 3-(2,4-dibenzoyloxyphenyl)-2-propenylaldehyde in the presence of NaB(OAc)3H in AcOH to give
 N-[N-[3-(2,4-dibenzoyloxyphenyl)-2-propenyl]-.beta.-O-benzyl-L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester, which was hydrogenated over Pd/C to afford I (R1 = R3 = OH, R2 = R4 = R5-R7 = H). The product tasted 10,000 times sweeter than sucrose.

L7 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2003 ACS
 TI N-Alkylaspartyldipeptide ester derivatives and sweeteners
 AN 2000:210206 CAPLUS
 DN 132:236239
 TI N-Alkylaspartyldipeptide ester derivatives and sweeteners
 IN Amino, Yusuke; Yuzawa, Kazuko; Takemoto, Tadashi; Nakamura, Ryoichiro
 PA Ajinomoto Co., Inc., Japan
 SO PCT Int. Appl., 44 pp.
 CODEN: PIXXD2

DT Patent.
 LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000017230	A1	20000330	WO 1999-JP4977	19990910
	W:	AU, BR, BY, CA, CN, CZ, HU, IL, IN, JP, KR, MX, NO, NZ, PL, RO, RU, SK, TR, UA, US, VN, ZA		JP 1998-264252	A 19980918
	RW:	AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		JP 1999-169419	A 19990616
CA 2344314	AA	20000330		CA 1999-2344314	19990910
				JP 1998-264252	A 19980918
				JP 1999-169419	A 19990616

AU 9956501	A1	20000410	WO 1999-JP4977 W 19990910
AU 748136	B2	20020530	AU 1999-56501 19990910
			JP 1998-264252 A 19980918
			JP 1999-169419 A 19990616
			WO 1999-JP4977 W 19990910
BR 9913779	A	20010710	BR 1999-13779 19990910
			JP 1998-264252 A 19980918
			JP 1999-169419 A 19990616
			WO 1999-JP4977 W 19990910
EP 1114828	A1	20010711	EP 1999-943302 19990910
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, SE, PT, IE, FI,			
RO			JP 1998-264252 A 19980918
			JP 1999-169419 A 19990616
			WO 1999-JP4977 W 19990910
NZ 510439	A	20020927	NZ 1999-510439 19990910
			JP 1998-264252 A 19980918
			JP 1999-169419 A 19990616
			WO 1999-JP4977 W 19990910
NO 2001001156	A	20010509	NO 2001-1156 20010307
			JP 1998-264252 A 19980918
			JP 1999-169419 A 19990616
			WO 1999-JP4977 W 19990910
US 2001039357	A1	20011108	US 2001-809197 20010316
			JP 1998-264252 A 19980918
			JP 1999-169419 A 19990616
			WO 1999-JP4977 W 19990910

OS MARPAT 132:236239

AB N-alkylaspartyldipeptide ester derivs. such as N-[N-[3-(3-hydroxy-4-methoxyphenyl)-3-methylbutyl]-L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester, are useful as sweeteners. The sweeteners have low caloric values and more sweet than conventional ones.

RE.CNT 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 10 OF 11 CAPLUS COPYRIGHT 2003 ACS
 TI Novel aspartyl dipeptide ester derivatives as sweeteners
 AN 2000:15228 CAPLUS
 DN 132:63481
 TI Novel aspartyl dipeptide ester derivatives as sweeteners
 IN Amino, Yusuke; Yuzawa, Kazuko; Takemoto, Tadashi; Nakamura, Ryoichiro
 PA Ajinomoto Co., Inc., Japan
 SO PCT Int. Appl., 28 pp.
 CODEN: PIXXD2
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000000508	A1	20000106	WO 1999-JP3050	19990607
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				

CA 2336133	AA	20000106	JP 1998-180204 A 19980626 CA 1999-2336133 19990607 JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607
AU 9940602	A1	20000117	AU 1999-40602 19990607
AU 752473	B2	20020919	JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607
EP 1088829	A1	20010404	EP 1999-923954 19990607
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, RO			JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607
BR 9911551	A	20011009	BR 1999-11551 19990607 JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607
NZ 508579	A	20021025	NZ 1999-508579 19990607 JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607
RU 2192430	C2	20021110	RU 2001-200110239919990607 JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607
NO 2000006627	A	20010212	NO 2000-6627 20001222 JP 1998-180204 A 19980626 WO 1999-JP3050 W 19990607

OS MARPAT 132:63481

AB The Markush structure of the aspartyl dipeptide ester derivs. (including salts thereof) are given, and the example is N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl]-L-(.alpha.-methyl)phenylalanine 1-Me ester. These compds. are low-calorie sweeteners and are sweeter than conventional ones.

RE.CNT 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 11 OF 11 CAPLUS COPYRIGHT 2003 ACS
TI Preparation of aspartyl dipeptide ester derivatives as sweeteners
AN 1999:672857 CAPLUS
DN 131:272186
TI Preparation of aspartyl dipeptide ester derivatives as sweeteners
IN Amino, Yusuke; Yuzawa, Kazuko; Takemoto, Tadashi; Nakamura, Ryoichiro
PA Ajinomoto Co., Inc., Japan
SO PCT Int. Appl., 36 pp.
CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9952937	A1	19991021	WO 1999-JP1210	19990311
W: AU, BR, BY, CA, CN, CZ, HU, IL, IN, JP, KR, MX, NO, NZ, PL, RO, RU, SK, TR, UA, US, VN				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD,				

TG

CA 2327938	AA	19991021	JP 1998-97701 A 19980409 JP 1999-38190 A 19990217 CA 1999-2327938 19990311 JP 1998-97701 A 19980409 JP 1999-38190 A 19990217
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AU 9941184	A1	19991101	WO 1999-JP1210 W 19990311
AU 753110	B2	20021010	AU 1999-41184 19990311
			JP 1998-97701 A 19980409
			JP 1999-38190 A 19990217
			WO 1999-JP1210 W 19990311
BR 9909542	A	20001226	BR 1999-9542 19990311
			JP 1998-97701 A 19980409
			JP 1999-38190 A 19990217
			WO 1999-JP1210 W 19990311
EP 1070726	A1	20010124	EP 1999-932431 19990311
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, RO			
			JP 1998-97701 A 19980409
			JP 1999-38190 A 19990217
			WO 1999-JP1210 W 19990311
RU 2179979	C1	20020227	RU 2000-200012801219990311
			JP 1998-97701 A 19980409
			JP 1999-38190 A 19990217
			WO 1999-JP1210 W 19990311
NZ 507938	A	20021126	NZ 1999-507938 19990311
			JP 1998-97701 A 19980409
			JP 1999-38190 A 19990217
			WO 1999-JP1210 W 19990311
ZA 9902566	A	19991012	ZA 1999-2566 19990407
			JP 1998-97701 A 19980409
NO 2000004979	A	20001107	NO 2000-4979 20001003
			JP 1998-97701 A 19980409
			JP 1999-38190 A 19990217
			WO 1999-JP1210 W 19990311

OS MARPAT 131:272186

AB Novel aspartyl dipeptide ester derivs. (including those in the form of a salt) having an excellent sweetening effect and usable as sweeteners such as N-[N-[3-(3-methyl-4-hydroxyphenyl)propyl]-L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester and

N-[N-[3-(3-hydroxy-4-methoxyphenyl)propyl]-L-.alpha.-aspartyl]-L-phenylalanine 1-Me ester (I) are prep'd. Thus, I was prep'd. from N-tert-butoxycarbonyl-.beta.-O-benzyl-.alpha.-L-aspartyl-L-phenylalanine Me ester and 3-benzyloxy-4-methoxycinnamaldehyde. I was 20,000-times sweeter than sucrose.

RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> logoff hold

COST IN U.S. DOLLARS

SINCE FILE ENTRY	TOTAL SESSION
---------------------	------------------

FULL ESTIMATED COST

35.34 244.84

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE ENTRY	TOTAL SESSION
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LOGINID: ssspta1623paz

PASSWORD :

* * * * * RECONNECTED TO STN INTERNATIONAL * * * * *
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FILE 'CAPLUS' ENTERED AT 07:39:28 ON 14 MAR 2003
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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	35.34	244.84
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-7.16	-9.11

=> logoff
ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y) / N / HOLD : Y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	35.34	244.84
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-7.16	-9.11

STN INTERNATIONAL LOGOFF AT 07:39:39 ON 14 MAR 2003

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: ssspta1623paz

STNLOGON timed out

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: ssspta1623paz

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS 2 Apr 08 "Ask CAS" for self-help around the clock
NEWS 3 Apr 09 BEILSTEIN: Reload and Implementation of a New Subject Area
NEWS 4 Apr 09 ZDB will be removed from STN
NEWS 5 Apr 19 US Patent Applications available in IFICDB, IFIPAT, and
IFIUDB
NEWS 6 Apr 22 Records from IP.com available in CAPLUS, HCAPLUS, and
ZCAPLUS
NEWS 7 Apr 22 BIOSIS Gene Names now available in TOXCENTER
NEWS 8 Apr 22 Federal Research in Progress (FEDRIP) now available
NEWS 9 Jun 03 New e-mail delivery for search results now available
NEWS 10 Jun 10 MEDLINE Reload
NEWS 11 Jun 10 PCTFULL has been reloaded
NEWS 12 Jul 02 FOREGE no longer contains STANDARDS file segment
NEWS 13 Jul 22 USAN to be reloaded July 28, 2002;
saved answer sets no longer valid
NEWS 14 Jul 29 Enhanced polymer searching in REGISTRY
NEWS 15 Jul 30 NETFIRST to be removed from STN
NEWS 16 Aug 08 CANCERLIT reload
NEWS 17 Aug 08 PHARMAMarketLetter(PHARMAML) - new on STN
NEWS 18 Aug 08 NTIS has been reloaded and enhanced
NEWS 19 Aug 19 Aquatic Toxicity Information Retrieval (AQUIRE)
now available on STN
NEWS 20 Aug 19 IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 21 Aug 19 The MEDLINE file segment of TOXCENTER has been reloaded
NEWS 22 Aug 26 Sequence searching in REGISTRY enhanced
NEWS 23 Sep 03 JAPIO has been reloaded and enhanced
NEWS 24 Sep 16 Experimental properties added to the REGISTRY file
NEWS 25 Sep 16 CA Section Thesaurus available in CAPLUS and CA
NEWS 26 Oct 01 CASREACT Enriched with Reactions from 1907 to 1985
NEWS 27 Oct 21 EVENTLINE has been reloaded
NEWS 28 Oct 24 BEILSTEIN adds new search fields
NEWS 29 Oct 24 Nutraceuticals International (NUTRACEUT) now available on
STN
NEWS 30 Oct 25 MEDLINE SDI run of October 8, 2002
NEWS 31 Nov 18 DKILIT has been renamed APOLLIT
NEWS 32 Nov 25 More calculated properties added to REGISTRY
NEWS 33 Dec 02 TIBKAT will be removed from STN
NEWS 34 Dec 04 CSA files on STN
NEWS 35 Dec 17 PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS 36 Dec 17 TOXCENTER enhanced with additional content
NEWS 37 Dec 17 Adis Clinical Trials Insight now available on STN
NEWS 38 Dec 30 ISMEC no longer available
NEWS 39 Jan 13 Indexing added to some pre-1967 records in CA/CAPLUS
NEWS 40 Jan 21 NUTRACEUT offering one free connect hour in February 2003
NEWS 41 Jan 21 PHARMAML offering one free connect hour in February 2003
NEWS 42 Jan 29 Simultaneous left and right truncation added to COMPENDEX,
ENERGY, INSPEC
NEWS 43 Feb 13 CANCERLIT is no longer being updated
NEWS 44 Feb 24 METADEX enhancements
NEWS 45 Feb 24 PCTGEN now available on STN
NEWS 46 Feb 24 TEMA now available on STN
NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation
NEWS 48 Feb 26 PCTFULL now contains images
NEWS 49 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results
NEWS 50 Mar 19 APOLLIT offering free connect time in April 2003

NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a,
CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),

NEWS HOURS AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002
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NEWS LOGIN General Internet Information
NEWS PHONE Welcome Banner and News Items
NEWS WWW Direct Dial and Telecommunication Network Access to STN
NEWS WWW CAS World Wide Web Site (general information)

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FILE COVERS 1907 - 20 Mar 2003 VOL 138 ISS 12
FILE LAST UPDATED: 19 Mar 2003 (20030319/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> aspartame derivative
2806 ASPARTAME
6 ASPARTAMES
2806 ASPARTAME
 (ASPARTAME OR ASPARTAMES)
40677 DERIVATIVE
299553 DERIVATIVES
338443 DERIVATIVE
 (DERIVATIVE OR DERIVATIVES)

547977 DERIV
912754 DERIVS
1238782 DERIV
(DERIV OR DERIVS)
1340774 DERIVATIVE
(DERIVATIVE OR DERIV)
L1 36 ASPARTAME DERIVATIVE
(ASPARTAME(W) DERIVATIVE)

=> solubility
53300 SOLUBILITY
19345 SOLUBILITIES
65619 SOLUBILITY
(SOLUBILITY OR SOLUBILITIES)
190561 SOLY
1 SOLIES
190561 SOLY
(SOLY OR SOLIES)
L2 207615 SOLUBILITY
(SOLUBILITY OR SOLY)

=> l1 and l2
L3 2 L1 AND L2

=> d l3 1-2 ti

L3 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS
TI Novel **aspartame derivative** crystal and process for
producing the same

L3 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS
TI Novel **aspartame derivative** crystal and process for
producing the same

=> aspartame
2806 ASPARTAME
6 ASPARTAMES
L4 2806 ASPARTAME
(ASPARTAME OR ASPARTAMES)

=> l4 and l2
L5 92 L4 AND L2

=> d 15 82-92 ti

L5 ANSWER 82 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI The physicochemical properties of .alpha.-sweet

L5 ANSWER 83 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI **Aspartame-mannitol resolidified fused mixture:** characterization
studies by differential scanning calorimetry, thermomicroscopy,
photomicrography and x-ray diffractometry

L5 ANSWER 84 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Dipeptide sweetener-metal complexes

L5 ANSWER 85 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Artificial sweetener tablets

L5 ANSWER 86 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Solubility improvement in dipeptide sweeteners

L5 ANSWER 87 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Aspartame administration to the infant monkey: hypothalamic morphology and plasma amino acid levels

L5 ANSWER 88 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Aspartame: a commercially feasible aspartic acid based sweetener

L5 ANSWER 89 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Concentrated liquid low calorie sweetener

L5 ANSWER 90 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Sweetener solution containing saccharin and aspartame

L5 ANSWER 91 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Easily dispersible, nonfoaming low-calorie sweetener

L5 ANSWER 92 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Cereal sweetening

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	13.35	13.56

SESSION WILL BE HELD FOR 60 MINUTES
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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	13.77	13.98

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FILE 'CAPLUS' ENTERED AT 14:14:15 ON 20 MAR 2003
L1 36 ASPARTAME DERIVATIVE
L2 207615 SOLUBILITY
L3 2 L1 AND L2
L4 2806 ASPARTAME

L5 92 L4 AND L2

=> d 15 71-81 ti

L5 ANSWER 71 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Manufacture of **aspartame**-containing beverages

L5 ANSWER 72 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Preparation of type 1 **aspartame** crystals with improved dissolution properties

L5 ANSWER 73 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Process for producing alpha-1-aspartyl-1-phenylalanine methyl ester having an improved **solubility**

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TI Applications of **aspartame** in soft drinks

L5 ANSWER 75 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Effect of sucrose, fructose and **aspartame** on fortificant iron **solubility** in a wheat flake cereal

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TI Incorporation of **aspartame** in sugar-containing food

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TI Readily soluble **aspartame** tablets

L5 ANSWER 78 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Readily soluble **aspartame** sweeteners

L5 ANSWER 79 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Artificially sweetened beverage mixes

L5 ANSWER 80 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Enhanced-**solubility aspartame** compounds

L5 ANSWER 81 OF 92 CAPLUS COPYRIGHT 2003 ACS
TI Stable dipeptide sweetener crystals and tablets containing the crystals

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COST IN U.S. DOLLARS
FULL ESTIMATED COST

SINCE FILE ENTRY	TOTAL SESSION
17.53	17.74

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